

Division of Environmental Health and Communicable Disease Prevention		
Section: 4.0 Diseases and Conditions	Updated 7/03	
Subsection: Lyme/Lyme-like Disease	Page 1 of 13	

Lyme/Lyme-like Disease Table of Contents

Lyme/Lyme-like Disease Fact Sheet CDC Lab Form (50.34) Lyme Disease Case Report (MO 580-1807)



Division of Environmental Health and Communicable Disease Prevention		
Section: 4.0 Diseases and Conditions	Updated 7/03	
Subsection: Lyme/Lyme-like Disease	Page 2 of 13	

Lyme/Lyme-like* Disease

*Lyme-like Disease: Missouri patients who fulfill the strict CDC surveillance definitions for Lyme disease are reported as such, although *Borrelia burgdorferi* has not been isolated from any of Missouri's cases, as of yet. The Missouri erythema migrans rashes are indistinguishable from those in other areas of the United States, but are referred to as erythema migrans-like by the Division of Vector-Borne Infectious Diseases, National Center for Infectious Diseases, Centers for Disease Control and Prevention. The clinical syndrome appears similar to Lyme disease. In Missouri, it is called Lyme-like disease. (1, 2)

Overview (3, 4)

For a complete description of Lyme disease, refer to the following texts:

- Control of Communicable Diseases Manual (CCDM).
- Red Book, Report of the Committee on Infectious Diseases.

Case Definition (5)

Clinical description

A systemic, tickborne disease with protean manifestations, including dermatologic, rheumatologic, neurologic, and cardiac abnormalities. The best clinical marker for the disease is the initial skin lesion (i.e., erythema migrans [EM]) that occurs in 60%-80% of patients.

Laboratory criteria for diagnosis

- Isolation of *Borrelia burgdorferi* from a clinical specimen or
- Demonstration of diagnostic immunoglobulin M or immunoglobulin G antibodies to *B. burgdorferi* in serum or cerebrospinal fluid (CSF). A two-test approach using a sensitive enzyme immunoassay or immunofluorescence antibody followed by Western blot is recommended.

Case classification

Confirmed: a) a case with EM or b) a case with at least one late manifestation (as defined below) that is laboratory confirmed.

Comment

This surveillance case definition was developed for national reporting of Lyme disease; it is not intended to be used in clinical diagnosis.



Division of Environmental Health and Communicable Disease Prever	
Section: 4.0 Diseases and Conditions	Updated 7/03
Subsection: Lyme/Lyme-like Disease	Page 3 of 13

Definition of terms used in the clinical description and case definition:

- Erythema migrans. For purposes of surveillance, EM is defined as a skin lesion that typically begins as a red macule or papule and expands over a period of days to weeks to form a large round lesion, often with partial central clearing. A single primary lesion must reach ≥5 cm in size. Secondary lesions also may occur. Annular erythematous lesions occurring within several hours of a tick bite represent hypersensitivity reactions and do not qualify as EM. For most patients, the expanding EM lesion is accompanied by other acute symptoms, particularly fatigue, fever, headache, mildly stiff neck, arthralgia, or myalgia. These symptoms are typically intermittent. The diagnosis of EM must be made by a physician. Laboratory confirmation is recommended for persons with no known exposure.
- Late manifestations. Late manifestations include any of the following when an alternate explanation is not found:
 - 1. *Musculoskeletal system*. Recurrent, brief attacks (weeks or months) of objective joint swelling in one or a few joints, sometimes followed by chronic arthritis in one or a few joints. Manifestations not considered as criteria for diagnosis include chronic progressive arthritis not preceded by brief attacks and chronic symmetrical polyarthritis. Additionally, arthralgia, myalgia, or fibromyalgia syndromes alone are not criteria for musculoskeletal involvement.
 - 2. *Nervous system*. Any of the following, alone or in combination: lymphocytic meningitis; cranial neuritis, particularly facial palsy (may be bilateral); radiculoneuropathy; or, rarely, encephalomyelitis. Encephalomyelitis must be confirmed by demonstration of antibody production against *B. burgdorferi in* the CSF, evidenced by a higher titer of antibody in CSF than in serum. Headache, fatigue, paresthesia, or mildly stiff neck alone are not criteria for neurologic involvement.
 - 3. *Cardiovascular system*. Acute onset of high-grade (2nd-degree or 3rd-degree) atrioventricular conduction defects that resolve in days to weeks and are sometimes associated with myocarditis. Palpitations, bradycardia, bundle branch block, or myocarditis alone are not criteria for cardiovascular involvement.
- Exposure. Exposure is defined as having been (≤30 days before onset of EM) in wooded, brushy, or grassy areas (i.e., potential tick habitats) in a county in which Lyme disease is endemic. A history of tick bite is not required.
- *Disease endemic to county.* A county in which Lyme disease is endemic is one in which at least two confirmed cases have been previously acquired or in which established populations of a known tick vector are infected with *B. burgdorferi*.



	able Disease Prevention	
	Section: 4.0 Diseases and Conditions	Updated 7/03
	Subsection: Lyme/Lyme-like Disease	Page 4 of 13

Information Needed for Investigation

Verify the diagnosis. Was there an EM rash of ≥ 5 cm? Determine what laboratory tests were conducted and the results.

Establish the extent of illness. Are other persons (e.g., family, friends) who might have shared the same exposure environment showing any signs of disease?

Case/Contact Follow Up And Control Measures

This disease is not transmitted person-to-person, so no contact follow up is required.

Control Measures

See the Control of Communicable Diseases Manual, Lyme Disease, "Methods of control."

See the <u>Red Book</u>, Lyme Disease, "Control Measures."

- Antibiotic prophylaxis after a tick bite (in the absence of symptoms) is usually not recommended. For this to be cost effective, the probability of *B. burgdorferi* infection would have to be abnormally high, which it is not in Missouri.
- As of February 25, 2002 the manufacturer announced that the LYMErixTM Lyme disease vaccine would no longer be commercially available. The efficacy of this vaccine against the Lyme-like disease seen in Missouri was never substantiated because the agent has not been positively identified and the vaccine had not been tested for efficacy against this disease.

Laboratory Procedures

The Missouri State Public Health Laboratory (SPHL) does not perform any testing for *B. burgdorferi*. However, the SPHL can arrange for testing of clinical specimens at various other laboratories. Information on laboratory procedures can be obtained from the Regional Communicable Disease Coordinator or from staff at the SPHL. The SPHL telephone number is 573-751-0633 and the web site is: http://www.dhss.state.mo.us/Lab/index.htm. (4 June 2003)

- Culture: *B. burgdorferi* organisms have been readily isolated from the margins of erythema migrans lesions; however, the organisms have been isolated infrequently from blood, joints, and cerebrospinal fluid. (6) Contact the SPHL for assistance in locating a laboratory that will isolate *B. burgdorferi*.
- **Serology:** The SPHL can submit serological specimens for testing to the Centers for Disease Control and Prevention, Fort Collins, Colorado. This laboratory offers an IgM



Division of Environmental Health and Communicable Disease Pr		
Section: 4.0 Diseases and Conditions	Updated 7/03	
Subsection: Lyme/Lyme-like Disease	Page 5 of 13	

antibody test on acute serum and an IgG antibody titer on paired serum. Positive or equivocal tests will be followed with a Western blot.

- CDC will perform IgM antibody testing on acute serum if the serum is collected within four weeks of onset. For best results the acute serum should be collected three to four weeks after onset of symptoms. Test results on specimens collected before three weeks may give a false negative result. A positive IgM test is indicative of a recent infection with *B. burgdorferi*.
- > CDC will perform IgG antibody testing on paired serum. For best results the acute serum should be collected three weeks after onset and the convalescent serum should be collected five weeks after the acute specimen was collected. A significant rise in IgG antibody titer between the acute and the convalescent sera is indicative of a recent infection with *B. burgdorferi*.
- All acute serum samples collected within the appropriate time frame will be sent to CDC for IgM testing and a request will be made at this time by the SPHL, to the submitter, for a convalescent specimen. Results of the IgM test will be sent to the submitter upon completion of testing. When the convalescent specimen arrives it will be sent to CDC for IgG testing. CDC will test both the acute and the convalescent specimens for IgG antibody. Upon completion of the IgG testing results will be sent to the submitter.
- ➤ <u>Submission of Blood Specimens</u>: The following forms must accompany specimens sent to the SPHL:
 - CDC Lab Form 50.34; in this Section or at: http://www.cdc.gov/ncidod/dvbid/misc/CDC50 34.pdf (4 June 2003)
 - 2. Syphilis test results by the RPR or MHA-TP test or
 - 3. A completed syphilis test request form, if the specimen is to be tested by the SPHL (this is to rule out syphilis, which is a spirochete, as is *B. burgdorferi*).
 - 4. A completed Lyme Disease Case Report (MO580-1807)

Reporting Requirements

Lyme disease is a Category II disease and shall be reported to the local health authority or to the Missouri Department of Health and Senior Services (DHSS) within three days of first knowledge or suspicion by telephone, facsimile or other rapid communication.

- 1. For all cases, complete a "Disease Case Report" (CD-1).
- 2. For all cases, complete a "Lyme Disease Case Report" (MO580-1807).
- 3. Entry of the completed CD-1 into MOHSIS negates the need for the paper CD-1 to be forwarded to the Regional Health Office.
- 4. Send the completed secondary investigation form to the Regional Health Office.



Division of Environmental Health and Communicable Disease Prevention		
Section: 4.0 Diseases and Conditions	Updated 7/03	
Subsection: Lyme/Lyme-like Disease	Page 6 of 13	

- 5. All outbreaks or "suspected" outbreaks must be reported as soon as possible (by phone, fax, or e-mail) to the Regional Communicable Disease Coordinator. This can be accomplished by completing the Missouri Outbreak Surveillance Report (CD-51).
- 6. Within 90 days of the conclusion of an outbreak, submit the final outbreak report to the Regional Communicable Disease Coordinator.

References

- 1. Lyme and/or Lyme-like Disease in Missouri. Masters, E.J., Donnell, H.D. Missouri Medicine. Vol 92, No. 7; July, 1995: 346-353.
- 2. Epidemiologic and Diagnostic Studies of Patients with Suspected Early Lyme Disease, Missouri, 1990-1993; Campbell, G.L., Paul, W.S., Schriefer, M.E., Craven, R.B., Robbins, K.E., Dennis, D.T. Journal of Infectious Diseases, 1995; 175:470-480.
- 3. Chin, James, ed. "Lyme Disease (Lyme borreliosis, Tickborne meningopolyneuritis)." <u>Control of Communicable Diseases Manual</u>. 17th ed. Washington, DC: American Public Health Association, 2000: 302-306.
- 4. American Academy of Pediatrics. "Lyme Disease (*Borrelia burgdorferi*)." In: Pickering, LK. ed. <u>2000 Red Book: Report of the Committee on Infectious Diseases</u>. 25th ed. Elk Grove Village, IL. 2000: 374-379.
- 5. Centers for Disease Control and Prevention. <u>Case Definitions for Infectious Conditions Under Public Health Surveillance</u>. MMWR 1997: 46 (No. RR-10). "Lyme Disease (*Borrelia burgdorferi*)," 1996, http://www.cdc.gov/epo/dphsi/casedef/lyme_disease_current.htm (4 June 2003)
- Moore, Kristine A, Hedberg, Craig, Osterholm, Michael T. "Lyme Disease." <u>Bacterial Infections of Humans Epidemiology and Control</u>. 3rd ed. Eds. Alfred S. Evans and Philip S. Brachman. New York: Plenum, 1998: 439-441.

Other Sources of Information

- 1. Steere, Allen C. "Borrelia Burgdorferi (Lyme Disease, Lyme Borreliosis)." <u>Principles and Practice of Infectious Diseases</u>. 5th ed. Eds. Gerald L. Mandell, John E. Bennett, and Raphael Dolin. New York: Churchill Livingstone, 2000:2504-2514.
- 2. Laboratory Serodiagnosis of Lyme Borreliosis: Callister, S.M.; Schell, R.F.; Journal of Spirochetal and Tick-borne Diseases; Volume 5, Spring/Summer 1998: 4-10.
- 3. <u>The Merck Veterinary Manual</u>. 8th Ed. Ed. Susan E. Aiello. Whitehouse Station, NJ: Merck & Co., Inc., 1998: 436, 2162. http://www.merckvetmanual.com/mvm/index.jsp (search "Lyme"). (4 June 2003)



Division of Environmental Health and Communicable Disease Prevention		
Section: 4.0 Diseases and Conditions	Updated 7/03	
Subsection: Lyme/Lyme-like Disease	Page 7 of 13	

Web Sites

- Centers for Disease Control and Prevention Protect Yourself from Ticks and Lyme Disease, http://www.cdc.gov/nasd/docs/d000901-d001000/d000961/d000961.html (4 June 2003)
- 2. Centers for Disease Control and Prevention Lyme Disease Home page, http://www.cdc.gov/ncidod/dvbid/lyme/index.htm (4 June 2003)
- 3. Lyme Disease Foundation, Inc., http://www.lyme.org (4 June 2003)
- 4. American Lyme Disease Foundation, Inc., http://www.aldf.com (4 June 2003)

Lyme/Lyme-like Disease

FACT SHEET

What is Lyme disease?

Lyme disease is an infectious illness that can be transmitted by the bite of a tick. It is caused by a type of bacteria called *Borrelia burgdorferi*.

What is Lyme-like disease?

Missouri patients who fulfill the strict Centers for Disease Control and Prevention (CDC) surveillance definition for Lyme disease are reported as such, although *B. burgdorferi* has not been isolated from any of Missouri's cases, as of yet. The Missouri erythema migrans rashes are indistinguishable from those in other areas of the United States, but are referred to as erythema migrans-like by CDC. The clinical syndrome appears similar to Lyme disease. In Missouri, it is called Lyme-like disease.

What are the symptoms of Lyme disease?

The symptoms of Lyme disease vary a great deal from one case to the next. In general, the early signs appear 3 to 32 days after a tick bite. The early symptoms of the disease can include fever, fatigue, headaches, aching joints, nausea, and a characteristic skin rash called erythema migrans (EM). EM, which occurs in 60-80 percent of the cases of Lyme disease, is roughly circular in shape. EM is usually found at the site of the tick bite, although it can also be found on other parts of the body. In time, the rash gets larger and the center often becomes clear (not reddened).

When the early symptoms are present it is important to treat the disease, otherwise late symptoms and complications can develop.

The late symptoms of the disease can include: severe headaches, stiff neck, weakness and/or pain in the extremities, facial paralysis, cardiac problems and arthritis (very common in this stage of the disease).

The late symptoms may occur weeks to years after being infected.

How does a person get Lyme disease?

The disease is transmitted through the bite of a tick infected with *B*. burgdorferi. The tick must be actually attached to a person's skin to transmit the infection. In experimental animals, transmission of Lyme disease does not occur until the tick has been attached for 24 hours or more; this may also be true in humans. Only a small percentage of ticks carry Lyme disease. The tick commonly known as the "deer tick" can carry the disease. Some other kinds of ticks may also spread Lyme disease, but this is not known with certainty. Because the tick may be very small, many people may not be aware that they have been bitten.

Can Lyme disease be treated?

Yes, once Lyme disease is diagnosed a physician will prescribe an antibiotic. Prompt treatment can cure the infection and usually prevents later complications.

Is there a vaccine for Lyme disease?

As of February 25, 2002 the manufacturer announced that the LYMErix™ Lyme disease vaccine would no longer be commercially available. The efficacy of this vaccine against the Lyme-like disease seen in Missouri was never substantiated because the agent has not been positively identified in Missouri and the vaccine had not been tested for efficacy against this disease.

How should a tick be removed?

Ticks should be removed promptly and carefully by using tweezers and applying gentle, steady traction. Do not crush the tick's body when removing it and apply the tweezers as close to the skin as possible to avoid leaving tick mouthparts in the skin. Do not remove ticks with your bare hands. Protect your hands with gloves, cloth, or tissue and be sure to wash your hands after removing a tick. After removing the tick, disinfect the skin with soap and water or other available disinfectants.

How can Lyme disease be prevented?

- 1. Avoid tick-infested areas, especially during the warmer months.
- 2. Wear light colored clothing so ticks can be easily seen and removed. Wear a long sleeved shirt, hat, long pants, and tuck your pant legs into your socks.
- 3. Walk in the center of trails to avoid overhanging grass and brush.
- 4. Check your body every few hours for ticks when you spend a lot of time outdoors in tick-infested areas. Ticks are most often found on the thigh, arms, underarms, legs or where tight fitting clothing has been.
- 5. Use insect repellents containing DEET on your skin or permethrin on your clothing. Be sure to follow the directions on the container and wash off repellents when going indoors. Carefully read the manufacturer's label on repellents before using on children.
- 6. Remove attached ticks immediately.

Missouri Department of Health and Senior Services Section for Communicable Disease Prevention Phone: (866) 628-9891 or (573) 751-6113

LABORATORY EXAMINATION(S) REQUESTED:			CATEGORY OF AGENT SUSPECTED:			
□ ANtimicrobial □ ISolation Susceptibility □ SErology (Specific Test) □ HIstology		☐ BActerial ☐ Rickettsial				
☐ HIstology ☐ SErology (Specific	Test)			■ PArasitic		
☐ IDentification ☐ OTher (Specify)			☐ FU ngal	OTher (Specify)		
SPECIFIC AGENT SUSPECTED:	OTHER ORGANISM(S) FOUND:	ISOLATION ATTEMPTED?	NO. OF TIMES	NO. OF TIMES	SPECIMEN SU	
			ISOLATED:	PASSED:	☐ Original Ma	aterial Mixed Isolate
		YES NO			Pure Isolate	e
DATE SPECIMEN TAKEN:	ORIGIN:			☐ OT her		
	<u> </u>			_		
SOURCE OF SPECIMEN:			SUBMITTED ON	J.		
	(Site)					
	e (Site)					
SErum SKin _						
SPutum STOOI	Specify)					
☐ UR ine ☐ TH roat ☐ OT her (S	pecify)		│ │ EGg │ (OTher (Specify)		
SERUM INFORMATION: MO DA YE	MO DA YR		IS AND SYMPTON	MS:		VOUS SYSTEM:
	□ 53/		FEver Maximum Temper	rature:	☐ HEadache ☐ MEningismu	IIS.
☐ ACute //			Duration:		■ MIcrocepha	alus
			CHills	Days	☐ HYdrocepha	alus
IMMUNIZATIONS:	МО	SKIN	l:		CErebral Ca	alcification
			MAculopapular		☐ CHorea	
(2.)			HEmorrhagic VEsicular		☐ PA ralysis ☐ OT her	
(3.)		🗖	Erythema Nodosu			
(4.)			Erythema Margina OTher	tum	MISCELLANEO ☐ JAundice	US:
TREATMENT: DRUGS USED None	DATE BEGUN DATE COM	PLETED			MYalgia	
	MO DA YR MO DA	YR RES	PIRATORY: RHinitis		☐ PLeurodyni☐ COnjunctivi	
		I <u>-</u>	PU lmonary		☐ CH orioretin	itis
(2.)		📙	PHaryngitis CAlcifications		☐ SPlenomeg ☐ HEpatomeg	•
(3.)		_	Otitis Media		Liver Absce	ess/cyst
EPIDEMIOLOGICAL DATA:			PNeumonia (type)		LYmphader	nopathy embrane Lesions
☐ SIngle Case ☐ SPoradic ☐ C	COntact		OTher		OTher	andrane Lesions
Family Illness			DIOVASCULAR:		STATE OF ILLN	Ecc.
Community Illness] MYocarditis] PEricarditis		☐ SYmptomat	
Travel and Residence (Location)			☐ ENdocarditis ☐ ASymptomatic ☐ OTher ☐ SUbacute		atic	
☐ Foreign		OTher		☐ CHronic		
_			TROINTESTINAL:		☐ DI sseminate	ed
_			Dlarrhea □ LOcalized □ BLood □ EXtraintestinal		inal	
			MUcous OTher			
Anthropod Contacts: None	Exposuer Only		COnstipation ABnormal Pain			
Type of Anthropod:		🗖	☐ VO miting			
Suspected Source of Infection:		U	OTher			
PREVIOUS LABORATORY RESULTS/0	THER CLINICAL INFORMATION: (Information)	ation supplied should	be related to this o	case and/or specime	en(s) and relative	to the test(s) requested.
			114117	FV	NUMBER	OUE
CDC 50.34 Rev. 09/2002 (BACK)	- CDC SPECIMEN SUBMISSION FORM	- CDC NU	UNIT MBER	FY	NUMBER	SUF.

Justification must be completed by State health department laboratory before specimen can be accepted by CDC. Please check the first applicable statement and when appropriate complete the statement with the ★. 1. Disease suspected to be of public health importance. Specimen is: (a) ☐ from an outbreak. (b) ☐ from uncommon or exotic disease. (c) ☐ an isolate that cannot be identified, is atypical, shows multiple antibiotic resistance, or from a normally sterile site(s) (d) ☐ from a disease for which reliable diagnostic reagents or expertise are unavailable in State. 2. ☐ Ongoing collaborative CDC/State project. Completed by:		STATE HEALTH DEPARTMENT LABORATORY ADDRE	SS:
*Prior arrangement for testing has been made. Please bring to the attention of:		STATE HEALTH DEPT. NO.:	DATE SENT TO CDC:
(Name):	Date://	PATIENT IDENTIFICATION: (Hospital No.)	(MM/DD/YYYY)/
Name, Address and Phone Number of Physician or O	rganization:	NAME:	
		(LAST, FIRST, MI) BIRTHDATE:	
		(MM/DD/YYYY)/	SEX: MALE FEMALE
		DIAGNOSIS:	
(FOR CDC USE ONLY) CDC NUMBER UNIT FY NUMBER	SUF DATE RECEIVED NO DA YR	ASSOCIATED ILLNESS:	
		DATE OF ONSET: (MM/DD/YYYY) ORM MUST BE COMPLETED	FATAL? YES NO
PLEA		THER PRINTED OR TYPED TE FORM FOR EACH SPECIMEN DATE REPORTED	
		MO DA YR	
	0 3	/	
	Comments:		
		D 6 5	
SERVICES OF SERVICES	Public He Centers for I	TH AND HUMAN SERVICES ealth Service Disease Control ectious Diseases	OC



nter for Infectious Diseas Atlanta, Georgia 30333



The Centers for Disease Control (CDC), an agency of the Department of Health and Human Services, is authorized to collect this information, including the Social Security number (if applicable), under provisions of the Public Health Service Act, Section 301 (42 U.S.C. 241). Supplying the information is voluntary and there is no penalty for not providing it. The data will be used to increase understanding of disease patterns, develop prevention and control programs, and communicate new knowledge to the health community. Data will become part of CDC Privacy Act system 09-20-0106, "Specimen Handling for Testing and Related Data" and may be disclosed: to appropriate State or local public health departments and cooperating medical authorities to deal with conditions of public health significance; to private contractors assisting CDC in analyzing and refining records; to researchers under certain limited circumstances to conduct further investigations; to organizations to carry out audits and reviews on behalf of HHS; to the Department of Justice in the event of litigation, and to a congressional office assisting individuals in obtaining their records. An accounting of the disclosures that have been made by CDC will be made available to the subject individual upon request. Except for permissible disclosures expressly authorized by the Privacy Act, no other disclosure may be made without the subject individual's written consent.



MO 580-1807 (6-93)

MISSOURI DEPARTMENT OF HEALTH BUREAU OF COMMUNICABLE DISEASE CONTROL

LYME DISEASE CASE REPORT	(CASE ID
NAME (LAST, FIRST, MIDDLE INITIAL)		TELEPHONE NUMBER
ADDRESS (STREET, CITY, STATE, ZIP CODE)		
COUNTY	PARENT'S NAME IF MINOR	
Mo Day Year	☐ Amer. Indian/Eskimo ☐ Asian/Pacific Isl. ☐ Black ☐ White ☐ Unknown Date of Diagnosis ? ☐ [Ethnicity
SYMPTOMS AND SIGNS OF CURRENT EP	ISODE (PLEASE MARK EACH	I QUESTION)
Erythema migrans (EM) present [N] [Y] [?] Physician diagnosed? [N] [Y] [?] EM first seen?		
RHEUMATOLOGIC: Arthritis characterized by brief attacks of joint swelling? [N] [Y] [?] Swelling observed by physician? [N] [Y] [?] Location of swollen joint(s):		
NEUROLOGIC: Did patient have Bell's palsy or cranial neuritis observed by physician? [N] [Y] [?] Describe cranial neuritis: Did patient have radiculoneuropathy? [N] [Y] [?] Name limb(s) affected: Did patient have lymphocytic meningitis? [N] [Y] [?] Did patient have encephalitis/encephalomyelitis? [N] [Y] [?]		
CARDIOLOGIC: 2nd or 3rd degree atrioventricular block? [N] [Y] [?] EKG done? [N] [Y] [?] Palpitations? [N] [Y] Cardiac manifestations:		

OVER

CLINICAL:		
Other late clinical symptoms:		
Was patient assessed for: Multiple sclerosis [N] [Y] [?] Rheumatoid fact	or [N] [Y] [?] Syphillis [N] [Y] [?]	
Evaluation		
OTHER I	HISTORY	
Was patient hospitalized? [N] [Y] [?]		
Name of antibiotic(s) used this episode?	No. of davs used:	
Was patient pregnant? [N] [Y] [?]		
LABORAT	ORY DATA	
CSF tested for antibodies to B burgdorferi? [N] [Y] [?]		
Blood tested for antibodies to <i>B burgdorferi?</i> [N] [Y] [?]		
Antibody to B burgdorferi higher in CSF than Serum? [N]	Y] [?]	
Serum Date Test Results	Serum Test Method Normal Values	
Mo. Day Year	Less than:	
Serology 1	U ELISA IFA III III III III III III III III II	
Serology 2 Day Year Grant Gran	ELISA IFA III	
Date of Culture Day Year Culture Results:	Western Blot Other	
CULTURE SITE	OTHER TESTS (SPECIFY)	
RESULTS		
EXPOSUR	F HISTORY	
EXPOSURE HISTORY Mo. Day Year		
Was there a tick bite? [N] [Y] [?] Date of tick bite:		
Site where tick bite (or other exposure, if not tick bite) occurred:	\bigcirc	
County State		
Type of tick: Lone Star Tick Dog Tick		
(Amblyomma americanum) (Dermace Black-legged Tick Unknown	ntor variabilis)	
(Ixodes scapularis)		
Other		
Activity at time of tick bite or other exposure?	Front Back	
PHYSICIAN'S NAME	PERSON COMPLETING FORM (IF DIFFERENT FROM PHYSICIAN)	
ADDRESS	ADDRESS	
,	ADDITION	
TELEPHONE NUMBER	TELEPHONE NUMBER	
FOR CENTRAL OFFICE STAFF USE ONLY		
Case Status: Meets Case Definition Does Not N	Meet Case Definition Pending	
Case Status. — Meets Case Demittion — Does NOT N	Meet Case Definition	